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Celgene Corporation*

**UNITED STATES DISTRICT COURT  
DISTRICT OF NEW JERSEY**

**CELGENE CORPORATION,**

**Plaintiff,**

**v.**

**HETERO LABS LIMITED, HETERO  
LABS LIMITED UNIT-V, HETERO  
DRUGS LIMITED, HETERO USA, INC.,  
AUROBINDO PHARMA LIMITED,  
AUROBINDO PHARMA USA, INC.,  
AUROLIFE PHARMA LLC, EUGIA  
PHARMA SPECIALTIES LIMITED,  
APOTEX INC., APOTEX CORP.,  
MYLAN PHARMACEUTICALS, INC.,  
MYLAN INC., MYLAN, N.V.,  
BRECKENRIDGE PHARMACEUTICAL,  
INC., and TEVA PHARMACEUTICALS  
USA, INC.,**

**Defendants.**

**Civil Action No. 17-3387 (ES)(MAH)  
(consolidated)**

**(Filed Electronically)**

**CONSENT JUDGMENT**

Plaintiff Celgene Corporation (“Celgene”) and Defendants Hetero Labs Limited, Hetero Labs Limited Unit-V, Hetero Drugs Limited, and Hetero USA, Inc. (collectively, “Hetero”), four of the parties in the above-captioned action, hereby stipulate and consent to entry of judgment and an injunction in this action as follows:

IT IS this 19th day of August, 2021:

ORDERED, ADJUDGED, AND DECREED as follows:

1. This Court has jurisdiction over the subject matter of the above action and has personal jurisdiction over the parties for purposes of this action only, including as set forth below in Paragraph 6 of this Consent Judgment.

2. As used in this Consent Judgment, the term “Hetero ANDA Product” shall mean a drug product manufactured, imported, sold, offered for sale, marketed, or distributed pursuant to Abbreviated New Drug Application No. 210236 in or for the United States of America, including its territories, possessions, and the Commonwealth of Puerto Rico.

3. As used in this Consent Judgment, the term “Patents-in-Suit” shall mean U.S. Patent Nos. 8,198,262; 8,673,939; 8,735,428; 8,828,427; 9,993,467; 10,555,939; 10,093,647; 10,093,648; and 10,093,649.

4. Until expiration of the Patents-in-Suit, Hetero, including any of its successors and assigns, is enjoined from infringing the Patents-in-Suit, on its own part or through any third party on its behalf, by making, having made, using, selling, offering to sell, importing, or distributing of the Hetero ANDA Product in or for the United States of America, including its territories, possessions, and the Commonwealth of Puerto Rico, unless and to the extent otherwise specifically authorized by Celgene, and is further enjoined from assisting or cooperating with any third parties in connection with any infringement of the Patents-in-Suit by any such third parties in connection with making, having made, using, selling, offering to sell, importing, or distributing of any pomalidomide-containing drug product that references NDA 204026 in or for the United States of America, including its territories, possessions, and the

Commonwealth of Puerto Rico, unless and to the extent otherwise specifically authorized by Celgene.

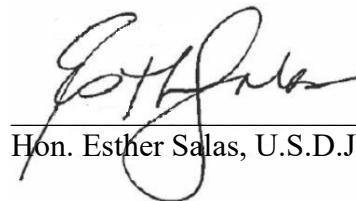
5. Compliance with this Consent Judgment may be enforced by Celgene and its respective successors in interest or assigns.

6. This Court retains jurisdiction to enforce the terms of this Consent Judgment and to enforce and resolve any disputes related thereto.

7. All claims, counterclaims, affirmative defenses and demands in this action that relate to Hetero are hereby dismissed with prejudice and without costs, disbursements, or attorneys' fees to any party.

8. Nothing herein prohibits or is intended to prohibit Hetero from maintaining any "Paragraph IV Certification" pursuant to 21 U.S.C. § 355(j)(2)(A)(vii)(IV) or pursuant to 21 C.F.R. § 314.94(a)(12) with respect to the Patents-in-Suit.

9. Nothing herein restricts or is intended to restrict the U.S. Food and Drug Administration from approving Abbreviated New Drug Application No. 210236 or the Hetero ANDA Product.



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Hon. Esther Salas, U.S.D.J.

We hereby consent to the form and entry of this Judgment:

Dated: August 18, 2021

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